LAMISIL AT- terbinafine hydrochloride aerosol, spray GlaxoSmithKline Consumer Healthcare Holdings (US) LLC

Drug Facts

Active ingredient

Terbinafine hydrochloride 1%

Purpose

Antifungal

Uses

- cures most jock itch (tinea cruris)
- relieves itching, burning, cracking, and scaling which accompany this condition

Warnings

For external use only

Do not use

- on nails or scalp
- in or near the mouth or the eyes
- for vaginal yeast infections

When using this product

- do not get into eyes. If contact occurs, rinse eyes thoroughly with water.
- do not puncture or incinerate; contents under pressure. Do not expose to heat or store at temperatures above 120°F (49°C).

Stop use and ask a doctor if

too much irritation occurs or gets worse

Keep out of reach of children.

If swallowed, get medical help or contact a Poison Control Center right away.

Directions

- adults and children 12 years and over:
 - wash the affected area with soap and water and dry completely before applying
 - to open remove clear cap
 - hold can 4" to 6" from skin. Press and hold to spray a thin layer over affected area
 - spray affected area once a day (morning **or** night) for 1 week or as directed by a doctor

- release to stop spray
- wipe excess from spray opening after each use
- return cap to can
- wash hands after each use
- children under 12 years: ask a doctor

Other information

store at 20° to 25° C (68° to 77° F)

Inactive ingredients

ethanol, polyoxyl 20 cetostearyl ether, propylene glycol, purified water

Questions?

call **1-800-330-9876**

Principal Display Panel

NDC 0067-6297-01

TERBINAFINE HYDROCHLORIDE SOLUTION 1% - ANTIFUNGAL

LAMISIL^{AT®}

SPRAY for JOCK ITCH

Cures most jock itch

For effective relief of itching and burning

Net Contents 125 ml (4.2 fl oz)

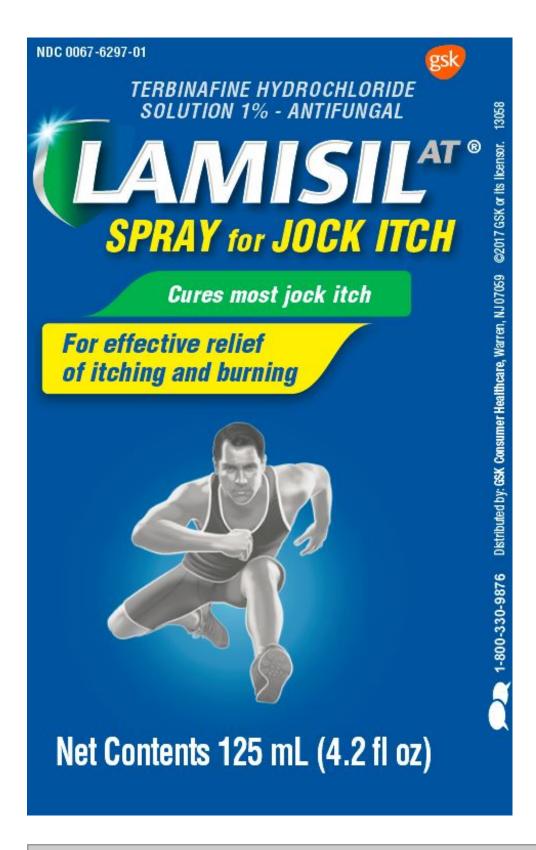
1-800-330-9876

Distributed by: **GSK Consumer Healthcare**, Warren, NJ 07059

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LAMISIL AT

terbinafine hydrochloride aerosol, spray

Product Information	Product Information				
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:0067-6297		
Route of Administration	TOPICAL				

Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
TERBINAFINE HYDRO CHLO RIDE (UNII: 012C11ZU6G) (TERBINAFINE - UNII:G7RIW8S0XP)	TERBINAFINE HYDROCHLORIDE	0.84 g in 125 mL	

Inactive Ingredients		
Ingredient Name	Strength	
ALCOHOL (UNII: 3K9958V90M)		
POLYOXYL 20 CETOSTEARYL ETHER (UNII: YRC528SWUY)		
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)		
WATER (UNII: 059QF0KO0R)		

ı	Packaging				
ı	#	Item Code	Package Description	Marketing Start Date	Marketing End Date
	1	NDC:0067-6297-01	125 mL in 1 CAN; Type 0: Not a Combination Product	03/07/2018	

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
NDA	NDA021124	03/07/2018		

Labeler - Glaxo SmithKline Consumer Healthcare Holdings (US) LLC (079944263)

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